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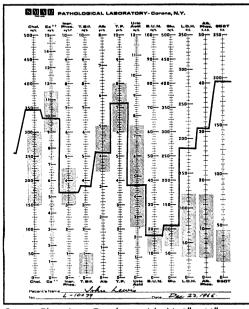
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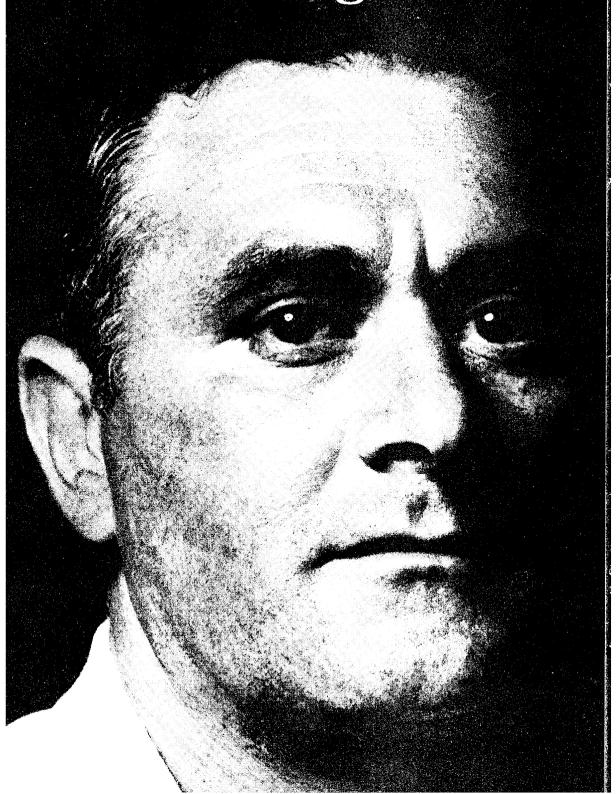
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The negative power of anxiety...

This man thinks he may never work again.



The patient who has had a myocardial infarction is usually advised by his physician to avoid emotional excitement. All too often his family, acutely concerned, transmits its anxiety to him, urging him to "rest, rest."

How anxiety may interfere: In a study of 336 males who had suffered at least one myocardial infarction, Sigler¹ reports that manual workers showed the lowest percentage of patients returning to work, compared to clerical workers, business and professional men. The author notes that in many cases the mere apprehension that "return to work would shorten life prevents the patient from resuming activities." It is also well known that emotional disturbance is probably the most common cause of cardiac disability in postinfarction cases.¹

The anxiety factor in both coronary and precoronary patients has recently been discussed by Thomas,² who suggests: "Intensive investigation of the sources and kinds of anxiety, and how destructive forms of anxiety can be identified and relieved may be the next important step in the prevention of coronary heart disease."

Relief of anxiety with Librium® (chlordiazepoxide HCl) often proves a valuable adjunct to medical counsel, reassurance and the total management program; may help prevent the postcoronary patient from regressing into a state of invalidism.

As an adjunct in cardiovascular therapy, Librium (chlordiazepoxide HCl): Quickly relieves anxiety of mild to severe degree in most cases. Helps expedite cooperation in therapeutic regimen. May be used concomitantly with certain specific medications of other classes of drugs, such as cardiac glycosides, antihypertensive agents and diuretics. By relieving anxiety, helps encourage productive activities. Has a wide margin of safety and, in proper maintenance dosage, seldom impairs mental acuity or ability to function. Often effective in extended therapy, usually without diminution of effect or need for increase in dosage—in protracted use, periodic blood counts and liver function tests are advisable.

References: 1. Sigler, L. H.: Geriatrics, 22:(9) 97, 1967. 2. Thomas, C. B.: Johns Hopkins Med. J., 122:69, 1968.

Before prescribing, please consult complete product information, a summary of which follows: Indications: Indicated when anxiety, tension and apprehension are significant components of the clinical profile. Contraindications: Patients with known hypersensitivity to the drug.

Warnings: Caution patients about possible combined effects with alcohol and other CNS depressants. As with all CNS-

acting drugs, caution patients against hazardous occupations requiring complete mental alertness (e.g., operating machinery, driving). Though physical and psychological dependence have rarely been reported on recommended doses, use caution in administering to addiction-prone individuals or those who might increase dosage; withdrawal symptoms (including convulsions), following discontinuation of the drug and similar to those seen with barbiturates, have been reported. Use of any drug in pregnancy, lactation, or in women of childbearing age requires that its potential benefits be weighed against its possible hazards. Precautions: In the elderly and debilitated, and in children over six, limit to smallest effective dosage (initially 10 mg or less per day) to preclude ataxia or oversedation, increasing gradually as needed and tolerated. Not recommended in children under six. Though generally not recommended, if combination therapy with other psychotropics seems indicated, carefully consider individual pharmacologic effects, particularly in use of potentiating drugs such as MAO inhibitors and phenothiazines. Observe usual precautions in presence of impaired renal or hepatic function. Paradoxical reactions (e.g., excitement, stimulation and acute rage) have been reported in psychiatric patients and hyperactive aggressive children. Employ usual precautions in treatment of anxiety states with evidence of impending depression; suicidal tendencies may be present and protective measures necessary. Variable effects on blood coagulation have been reported very rarely in patients receiving the drug and oral anticoagulants; causal relationship has not been established clinically. Adverse Reactions: Drowsiness, ataxia and confusion may occur, especially in the elderly and debilitated. These are reversible in most instances by proper dosage adjustment, but are also occasionally observed at the lower dosage ranges. In a few instances syncope has been reported. Also encountered are isolated instances of skin eruptions. edema, minor menstrual irregularities, nausea and constipation, extrapyramidal symptoms, increased and decreased libido - all infrequent and generally controlled with dosage reduction; changes in EEG patterns (lowvoltage fast activity) may appear during and after treatment; blood dyscrasias (including agranulocytosis), jaundice and hepatic dysfunction have been reported occasionally, making periodic blood counts and liver function tests advisable during protracted therapy.

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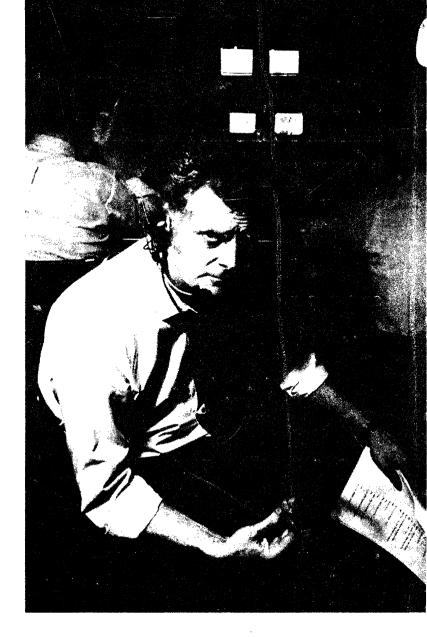
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Valium® (diazepam)

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Before prescribing, please consult complete product information, a summary of which follows: Indications: Tension and anxiety states; somatic complaints which are concomitants of emotional factors; psychoneurotic states manifested by tension, anxiety, apprehension, fatigue, depressive symptoms or agitation; acute agitation, tremor. delirium tremens and hallucinosis due to acute alcohol withdrawal; adjunctively in skeletal muscle spasm due to reflex spasm to local pathology, spasticity caused by upper motor neuron disorders, athetosis, stiff-man syndrome, convulsive disorders (not for sole therapy). Contraindicated: Known hypersensitivity to the drug. Children under 6 months of age. Acute narrow angle

Warnings: Not of value in psychotic patients. Caution against hazardous occupations requiring complete mental alertness. When used adiunctively in convulsive disorders,



possibility of increase in frequency and/or severity of grand mail seizures may require increased dosage of standard anticonvulsant medication; abrupt withdrawal may be associated with temporary increase in frequency and/or severity of seizures. Advise against simultaneous ingestion of alcohol and other CNS depressants. Withdrawal symptoms have occurred following abrupt discontinuance. Keep addiction-prone individuals under careful surveillance because of their predisposition to habituation and dependence. In pregnancy, lactation or women of childbearing age, weigh potential benefit against possible hazard.

Precautions: If combined with other psychotropics or anticonvulsants, consider carefully pharmacology of agents employed. Usual precautions indicated in patients severely depressed, or with latent depression, or with suicidal tendencies. Observe usual precautions in impaired renal or hepatic function. Limit dosage to

smallest effective amount in elderly and debilitated to preclude ataxia or oversedation.

Side Effects: Drowsiness, confusion, diplopia, hypotension, changes in libido, nausea, fatigue, depression, dysarthria, jaundice, skin rash, ataxia, constipation, headache, incontinence, changes in salivation. slurred speech, tremor, vertigo, urinary retention, blurred vision. Paradoxical reactions such as acute hyperexcited states, anxiety, hallucinations, increased muscle spasticity. insomnia, rage, sleep disturbances, stimulation, have been reported; should these occur, discontinue drug. Isolated reports of neutropenia, jaundice; periodic blood counts and liver function tests advisable during long-term therapy.

